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R² is an acyl group;

 CO_2R^3 is a carboxy group or a carboxylate anion, or R^3 is a readily removable carboxy protecting group;

R⁴ represents hydrogen or up to four substituents selected from alkyl, alkenyl, alkynyl, alkoxy, hydroxy, halogen, amino, alkylamino, acylamino, dialkylamino, CO₂R, CONR₂, SO₂NR₂ (where R is hydrogen or C₁₋₆ alkyl), aryl and heterocyclyl, which may be the same or different and wherein any R⁴ alkyl substituent is optionally substituted by any other R⁴ substituent;

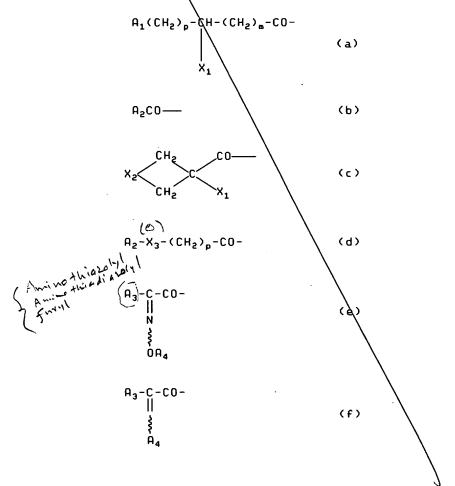
X is S, SO, SO₂, O or CH_2 ;

m is 1 or 2;

n is 0;

"acyl" is selected from the group consisting of formula

(a) to (f):



wherein p is 0, 1 or 2;

 $m \in 0, 1 \text{ or } 2;$

 A_1 is $C_{1.6}$ alkyl, substituted $C_{1.6}$ alkyl, $C_{3.6}$ cycloalkyl, cyclohexenyl, cyclohexadienyl, or an aromatic or heteroaromatic group;

X₁ is a hydrogen or halogen atom, a carboxylic acid, carboxylic ester, sulphonic acid, azido, tetrazolyl, hydroxy, acyloxy, amino, ureido, acylamino, heterocyclylamino, guanidino or acylureido group;

A₂ is an aromatic or-heteroaromatic group, a substituted alkyl group, or a substituted dithietane;

X₂ is a -CH₂OCH₂-, -CH₂SCH₂- or alkylene group;

X₃ is an oxygen or sulphur atom;

A₃ is an aryl or heteroaryl group; and

 A_4 is hydrogen, $C_{1.6}$ alkyl, $C_{3.8}$ cycloalkyl, $C_{3.8}$ cycloalkyl($C_{1.6}$) alkyl, $C_{1.6}$ alkoxycarbonyl($C_{1.6}$) alkyl, $C_{2.6}$ alkenyl, carboxy($C_{1.6}$) alkyl, $C_{2.6}$ alkynyl, aryl or $C_{1.6}$ alkyl substituted by up to three aryl groups.

56. A compound as claimed in claim 57 having the formula (Ia):

$$R^{2}NH$$
 R^{1}
 R^{1}
 $R^{2}NH$
 $R^$

wherein R^1 , R^2 , R^4 , m, n and X are as defined with respect to formula (I) in claim 57 and the group CO_2R^6 is CO_2R^3 where CO_2R^3 is a carboxy group or a carboxylate anion, or a pharmaceutically acceptable salt or in vivo hydrolysable ester thereof.

59? A compound as claimed in claim 57 wherein R^1 is hydrogen.

LA21952/35522

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optionally substituted phenyl, X_1 is hydrogen or amino, A_2 is optionally substituted phenyl, X_3 is oxygen, A_3 is aminothiazolyl, aminothiadiazolyl or furyl, and R^4 is hydrogen, $C_{1.6}$ alkyl, or carboxy $C_{1.6}$ alkyl./

A compound as claimed in claim 57 wherein CO_2R^3 is carboxy or a carboxylate anion or R^3 is \underline{t} -butyl, 4-methoxybenzyl, diphenylmethyl, acetoxymethyl, acetoxyethyl, pivaloyloxymethyl, propan-2-yloxycarbonyloxyethyl or 2-ethoxycarbonyl-but-2-enyl.

A compound as claimed in claim $\mathcal M$ wherein the cyclic ether group bonded to the 3-position of the cephalosporin nucleus is unsubstituted or substituted by up to three substituents selected from C_{1-6} alkyl, C_{1-6} alkoxy, C_{1-6} alkoxy carbonyl, C_{1-6} alkanoyloxy C_{1-6} alkyl or C_{1-6} alkoxy C_{1-6} alkyl. $\mathcal M$

A compound as claimed in claim 51 wherein m is 1.

A compound as claimed in claim 51 wherein the cyclic ether group is a tetrahydrofuran-2-yl or a tetrahydropyran-2-yl group.

68. A compound as claimed in claim 57 selected from the group consisting of:

sodium (6R,7R)-7-[2-(2-aminothiazol-4-yl)-2-(Z)-methoxy-iminoacetamido]-3-[(RS)-tetrahydrofuran-2-yl]ceph-3-em-4-carboxylate;

pivaloyloxymethyl (6R, 7R) -7-[2-(2-aminothiazol-4-yl)-2-(\underline{Z})-methoxyiminoacetamido]-3-[(\underline{RS})-tetrahydrofuran-2-yl]ceph-3-em-4-carboxylate;

sodium (6R,7R)-7-[2-(2-aminothiazol-4-yl)-2-(\underline{Z})-methoxy-iminoacetamido]-3-[(\underline{RS})-tetrahydropyran-2-yl]ceph-3-em-4-carboxylate;

pivaloyloxymethyl (6R, 7R)-7-[2-(2-aminothiazol-4-yl)-2-(\underline{Z})-methoxyiminoacetamido]-3-[(\underline{RS})-tetrahydropyran-2-yl]ceph-3-em-4-carboxylate;

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(6R, 7R) -7-[2-(2-aminothiazol-4-yl)-2-(Z)-hydroxy-iminoacetamido]-3-[(RS)-tetrahydrofuran-2-yl]ceph-3-em-4-carboxylic acid;

sodium (6R,7R)-7-[2-(2-aminothiazol-4-yl)-2-(Z)-methoxy-iminoacetamido]-3-[(S)-tetrahydrofuran-2-yl]ceph-3-em-4-carboxylate;

pivaloyloxymethyl $(6\underline{R}, 7\underline{R})$ -7-[2-(2-aminothiazol-4-yl)-2-(\underline{Z})-methoxyiminoacetamido]-3-[(\underline{S})-tetrahydrofuran-2-yl]ceph-3-em-4-carboxylate;

sodium (6R,7R)-7-[2-(2-aminothiazol-4-yl)-2-(Z)-methoxy-iminoacetamido]-3-[(R)-tetrahydrofuran-2-yl]ceph-3-em-4-carboxylate;

pivaloyloxymethyl $(6\underline{R}, 7\underline{R})$ -7-[2-(2-aminothiazol-4-yl)-2-(\underline{Z})-methoxyiminoacetamido]-3-[(\underline{R})-tetrahydrofuran-2-yl]ceph-3-em-4-carboxylate;

diphenylmethyl (6R, 7R)-7-phenylacetamido-3-[(RS)-tetrahydrofuran-2-yl]ceph-3-em-4-carboxylate;

sodium (6R,7R)-7-[2-(2-aminothiazol-4-yl)-2-(\underline{Z})-methoxy-iminoacetamido]-3-[(\underline{RS})-tetrahydrofuran-3-yl]ceph-3-em-4-carboxylate;

acetoxymethyl $(6\underline{R}, 7\underline{R})$ -7-[2-(2-aminothiazol-4-yl)-2-(\underline{Z})-methoxyiminoacetamido]-3-[(\underline{S})-tetrahydrofuran-2-yl]ceph-3-em-4-carboxylate;

sodium $(6\underline{R}, 7\underline{R})$ -7-[2-(2-aminothiazol-4-yl)-2-(\underline{Z})-methoxy-iminoacetamido]-3-(5-methoxymethyltetrahydrofuran-2-yl]ceph-3-em-4-carboxylate;

sodium (6R,7R)-7-[2-(2-aminothiazol-4-yl)-(Z)-pent-2-enamido]-3-[(S)-tetrahydrofuran-2-yl]ceph-3-em-4-carboxylate;

sodium $(6\underline{R}, 7\underline{R}) - 7 - [2 - (2 - aminothiazol - 4 - y1) - 2 - (\underline{Z}) - methoxyiminoacetamido] - 3 - [(\underline{S}) - tetrahydrofuran - 2 - y1]ceph - 3 - em - 4 - carboxylate;$

 $\frac{(RS)}{-1-acetoxyethyl} (6R,7R)-7-[2-(2-aminothiazol-4-yl)-2-(Z)-methoxyiminoacetamido]-3-[(S)-tetrahydrofuran-2-yl]ceph-3-em-4-carboxylate;$

(6R, 7R) - 7 - [2 - (2 - aminothiazol - 4 - y1) - 2 - (Z) - carboxy-methoxyiminoacetamido] - 3 - [(RS) - tetrahydrofuran - 2 - y1]ceph - 3 -

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em-4-carboxylic acid disodium salt;

sodium (6R, 7R)-7-[(R)-2-amino-2-(4-hydrophenyl)-acetamido]-3-[(S)-tetrahydrofuran-2-yl]ceph-3-em-4-carboxylate;

sodium $(1\underline{S}, 6\underline{R}, 7\underline{R})$ -7-[2-(2-aminothiazol-4-yl)-2-(\underline{Z})-methoxyiminoacetamido]-3-[(\underline{S})-tetrahydrofuran-2-yl]ceph-3-em-4-carboxylate-1-oxide;

sodium 7-[2-(2-aminothiazol-4-yl)-2-(<u>Z</u>)-methoxyimino-acetamido]-3-(tetrahydrofuran-2-yl)-1-carba-1-dethiaceph-3-em-4-carboxylate;

sodium $(6\underline{R}, 7\underline{R})$ -7-[2-(2-aminothiazol-4-yl)-2-(\underline{Z})-methoxyiminoacetamido]-3-[(\underline{S})-tetrahydrofuran-2-yl]ceph-3-em-4-carboxylate-1,1-dioxide;

(RS)-1-(propan-2-yl) oxycarbonyloxyethyl (6R, 7R)-7-[2-(2-aminothiazol-4-yl)-2-(\underline{Z})-methoxyiminoacetamido]-3-[(\underline{S})-tetrahydrofuran-2-yl]ceph-3-em-4-carboxylate;

sodium (6R, 7R)-7-[2-(2-aminothiazol-4-yl)-2-(\underline{Z})-methoxyiminoacetamido]-3-[(5R, 2SR)-5-methyltetrahydrofuran-2-yl]ceph-3-em-4-carboxylate;

sodium (6R,7R)-7-[2-(furan-2-yl)-2-(\underline{Z})-methoxyimino-acetamido]-3-[(\underline{S})-tetrahydrofuran-2-yl]ceph-3-em-4-carboxylate;

sodium $(6\underline{R}, 7\underline{R})$ -7-[2-(2-aminothiazol-4-yl)-2-(\underline{Z})-methoxy-iminoacetamido]-3-[(\underline{S})-5,5-dimethyltetrahydrofuran-2-yl]ceph-3-em-4-carboxylate;

sodium (6R, 7R)-7-[2-(2-aminothiazol-4-yl)-2-(\underline{Z})-methoxy-iminoacetamido]-3-(5)-methoxycarbonyltetrahydrofuran-2-yl)ceph-3-em-4-carboxylate;

sodium (6R, 7R)-7-[2-(2-aminothiazol-4-yl)-2-(\underline{Z})-methoxy-iminoacetamido]-3-[3-methyltetrahydrofuran-2-yl]ceph-3-em-4-carboxylate; and

2-ethoxycarbonyl- (\underline{Z}) -but-2-enyl $(6\underline{R},7\underline{R})$ -7- $[2-(2-aminothiazol-4-yl)-2-(\underline{Z})$ -methoxyiminoacetamido]-3- $[(\underline{S})$ -tetrahydrofuran-2-yl]ceph-3-em-4-carboxylate.

of claim 58 or a pharmaceutically acceptable salt or <u>in vivo</u>

LA21952/35522